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(21) International Application Number: PCT/NL99/00676 (22) International Filing Date: 4 November 1999 (04.11.99) (30) Priority Data: 98203719.4 4 November 1998 (04.11.98) EP (71) Applicant (for all designated States except US): ENGELHARD CORPORATION [US/US]; 101 Wood Avenue, Iselin, NJ 08830-0770 (US). (72) Inventors; and (75) Inventors/Applicants (for US only): BAYENSE, Cornelis, Roeland [NL/NL]; Delkant 6, NL-5311 CJ Gameren (NL). YKEMA, Durk [NL/NL]; Hemelvuur 13, NL-3454 SP De Meern (NL). (74) Agent: OTTEVANGERS, S., U.; Vereenigde, Nieuwe Parklaan 97, NL-2587 BN The Hague (NL).		(81) Designated States: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZW, ARIPO patent (GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG). Published <i>With international search report.</i>
(54) Title: STAR SHAPED ALUMINA EXTRUDATES AND CATALYST BASED THEREON (57) Abstract <p>This invention is directed to star shaped alumina extrudates with a pore volume in the pores of a diameter over 1000 nm, as determined by mercury porosity, of at least 0.05 ml/g and a total pore volume between 0.5-0.75 ml/g. The extrudates have a length of between 2-8 mm, a length to diameter ratio of between 1-3, a side crushing strength of at least 50 N and a bulk crushing strength of at least 1 MPa.</p>		

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Title: STAR SHAPED ALUMINA EXTRUDATES AND CATALYST BASED THEREON

The invention is directed to transition alumina extrudates, suitable as catalyst, or as catalyst support, and the use of such extrudates in chemical reactions.

In catalysis alumina plays an important role, both as
5 a catalyst support and as catalytically active material. As is mentioned in Kirk-Othmer, Third Ed, Vol. 2, pages 230-232, alumina can be used as catalyst in a Claus process, for dehydration of alcohols, such as the production of olefins from alcohol, and the reverse reaction, but also for the
10 isomerisation of olefins. As interacting catalyst support alumina may play a role in hydrorefining catalysts, e.g. in cobalt or nickel-molybdenum oxides on alumina.

As a support alumina is frequently used for precious metal catalyst, such as in exhaust catalysts or for
15 (de)hydrogenation reactions. As support for a nickel catalyst it may be used in (de)hydrogenation reactions such as for fat and oils hydrogenation, for hydrogenation of fatty nitriles or of nitro aromatic compounds or for oligomerisation of olefins.

20 The structure of the support, i.e. the BET surface area, the pore size and the pore volume distribution, forms an important aspect of the alumina or alumina based catalyst. In view of activity and selectivity it would be highly desirable to have an alumina product that is on the one hand
25 highly porous, i.e. having a large volume in large pores, and that has a good mechanical strength and stability. Unfortunately these are requirements that are difficult to reconcile with each other.

In fixed bed processes shaped bodies of alumina are
30 frequently used. An important aspect therein is the shape dependency of the pressure drop. Tablets and extrudates are the materials commonly used in fixed bed applications. In

order to minimise pressure drop, the use of star shaped extrudates would be most suitable. However, star shaped bodies, extrudates, tend to be prone to attrition due to the presence of the 'points' of the star.

5 It is an object of the invention to reconcile these various requirements in the form of a transition alumina extrudate, having a carefully balanced set of properties. Further objects and advantages will become clear from the following description of the invention and the preferred
10 embodiments thereof.

 The invention is based thereon that the inventors have now been able to provide a star shaped alumina extrudate, having on the one hand an optimal structure, as indicated above and on the other hand a good strength.

15 The invention is accordingly directed to star shaped alumina extrudates with a pore volume in pores of diameter of over 1000 nm, as determined by mercury porosimetry, of at least 0.05 ml/g, a side crushing strength of at least 50 N and a bulk crushing strength of at least 1 MPa.

20 Surprisingly, this set of properties can be made available in one material, thereby providing a material with which chemical reactions can be made much more efficient, resulting in higher activity and/or selectivity. Also the material of the invention, when used in fixed bed reactors,
25 provides a decreased pressure drop compared to regular extrudates having a cylindrical shape.

 The BET surface area, as determined by single point adsorption using the BET equation (as e.g. described by G. Sandstede et.al., Chem. Ing. Tech. 32 (1960), 413), should be
30 at least 10 m²/g of alumina. This coincides with the requirement of using a transition alumina, i.e. not an α alumina. Suitable alumina's are the various transition alumina's including γ -alumina, δ -alumina, ϵ -alumina, κ -alumina, ζ -alumina, θ -alumina and τ -alumina. These alumina's
35 have a large BET-surface area, generally in the range of 25 up to more than 100 m²/g.

The pore volume is a further important requirement, whereby it is on the one hand important that the total pore volume, as determined by mercury intrusion is sufficiently high and on the other hand that the pore volume in pores of over 1000 nm forms a substantial portion of the total pore volume. In absolute terms the total pore volume should be at least 0.50 ml/g, whereas the ratio of the pore volume in pores of over 1000 nm to total pore volume should preferably be more than 0.04. An alumina having those properties has good properties in terms of reactant accessibility, which makes it very suitable for all kinds of catalytic reactions requiring good diffusion of reactants and products through the alumina, thereby eliminating diffusion limitation problems as much as possible.

The pore volume and pore size distribution are determined by mercury porosimetry measurements, as described by J. Rouquerol et al in Pure & Applied Chem., 66(8),1994, pages 1752-1753, using the Washburn equation.

As indicated above, the use of star shaped extrudates is important in terms of pressure drop in relation to accessibility of the internal surface of the alumina. This also plays a role in eliminating diffusion problems. Star shaped extrudates can be defined as objects having some kind of central part or core, with three or more triangularly shaped extensions on the circumference thereof. Most preferred are star shaped extrusions having five extensions, as this provides the optimal balance between strength, porosity, pressure drop and accessibility. Another advantageous property of the star shaped extrudates is the fact that the ratio of external surface area to volume is more advantageous than in the case of conventional cylindrical extrudates or tablets.

The ratio of the length of the extrudates to the diameter is preferably between 1 and 3, whereby as diameter the distance is meant between two parallel planes on either side of the extrudate.

Important aspects of the material of the invention are also the strength characteristics. As indicated above a side crushing strength of at least 50 N and a bulk crushing strength of at least 1 MPa are essential herein. These
5 parameters form the basis for the suitability of the extrudates for use in large scale reactors, like in the petroleum industry. When the extrudates meet these requirements, they can be used in huge fixed bed reactors, that require very strong material. The side crushing strength
10 and the bulk crushing strength is defined as follows:

The side crushing strength (SCS) of extrudates is defined as the pressure (in Newtons) at which extrudates of 4.5-5.00 mm length are crushed, when treated under pressure between two flat plates on a AIKOH, 9500 series tester.

15 The bulk crushing strength (BCS) of a catalyst is defined as the pressure (in Megapascals) at which 0.5% fines (i.e. particles less than 0.425 mm) are formed when treated under a piston in a tube. For that purpose, 17 ml of catalyst particles, presieved on a 0.425 mm sieve, are loaded in a
20 cylindrical sample tube (diameter 27.3 mm), and 8 ml steel beads is loaded on top. The catalyst is subsequently treated at different (increasing) pressures for three minutes, after which the fines are recovered and their percentages is determined. This procedure is repeated until a level of
25 0.5 wt.% fines is reached.

Another aspect of the strength of the material is the attrition, i.e. the amount of material that may break off of the extrudates upon use. This attrition, determined in accordance with ASTM D4058-87, should preferably be less than
30 5 wt.%, more in particular less than 3 wt.%.

The alumina extrudates having the above properties can be prepared by mixing transition alumina powder with a suitable binder in the presence of a liquid, usually water or an aqueous solution of a mineral acid such as hydrochloric,
35 sulfonic or nitric acid, to form a paste, followed by extruding of the paste in the required star form, using a

suitable die and cutting the extruded strands of material to the required length. Optionally after drying, the extrudates are calcined.

It is possible to use various types of binder materials, such as those based on silica or alumina. Examples are colloidal silica, waterglass, or clays. It is preferred to use an alumina based binder or a binder that is removed during calcination, while providing and maintaining the required strength. An example of a suitable binder system is an alumina binder that gels under acidic treatment, for example by using organic or inorganic acids. The amount of binder material used in the preparation of the paste that is to be extruded will vary depending on the type of material and the required strength. Generally it will not be in excess of 30 wt.% based on the dry weight of binder and alumina together.

The invention will now be elucidated on the basis of an example.

EXAMPLE

20

1.5kg of aluminium trihydrate, containing 65 wt.% of Al_2O_3 , with an average particle size of 30-50 μm is mixed with 0.4 kg of alumina binder. The powders are mixed extensively while slowly adding diluted, aqueous HNO_3 in an amount of 2 wt.%, calculated on the weight of the total amount of alumina.

Thereby the alumina binder is peptised. Mixing is continued until a relatively dry product is obtained. the intermediate product is extruded using a one-screw extruder, equipped with a die having starshaped holes and a cutting device.

The extrudates obtained are dried at 105°C for 16 hours and subsequently calcined at 850°C for one hour. Attached are two figures with photographs of an extrudate shown from two different angles.

The final product has been analysed for its physical properties with the following result:

N₂-BET surface area 106 m/g²

Total Hg pore volume 0.56 ml/g

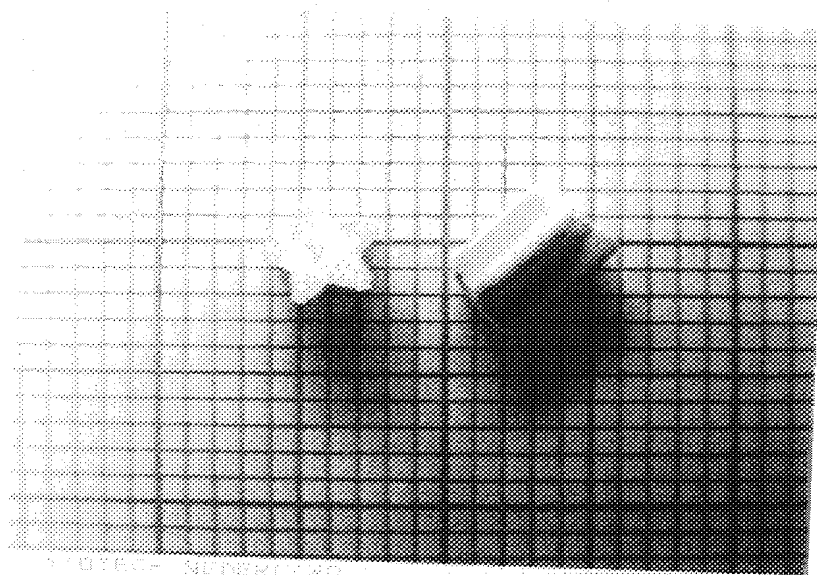
5 Pore volume in pores over 1000 nm 0.07 ml/g

Side crushing strength 65 N

Bulk crushing strength 1.08 MPa

Claims

1. Star shaped alumina extrudates with a pore volume in pores of diameter of over 1000 nm, as determined by mercury porosimetry, of at least 0.05 ml/g, a side crushing strength of at least 50 N and a bulk crushing strength of at least
5 1 MPa.
2. Extrudates according to claim 1, having a length of between 2 and 8mm.
3. Extrudates according to claim 1 or 2, having a length to diameter ratio of between 1 and 3.
- 10 4. Extrudates according to claims 1-3, wherein the total pore volume a determined by mercury porosimetry is between 0.5 and 0.75 ml/g.
5. Extrudates according to claims 1-4, wherein the BET surface area is at least 75 m²/g.
- 15 6. Extrudates according to claims 1-5, wherein the attrition in accordance with ASTM D4058-87 is less than 5 wt.%, preferably less than 3 wt.%.
7. Catalyst, comprising at least one catalytically active material supported on an extrudate according to claims
20 1-6.
8. Catalyst according to claim 7, wherein the catalytically active material is selected from the group of metals, metal oxides, metal sulfides and combinations thereof.
- 25 9. Use of an extrudate according to claims 1-6 or a catalyst according to claim 7 or 8 in a chemical reaction.



INTERNATIONAL SEARCH REPORT

International Application No

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A. CLASSIFICATION OF SUBJECT MATTER
IPC 7 B01J35/02 B01J21/04

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 B01J

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	EP 0 008 424 A (BASF AG) 5 March 1980 (1980-03-05) claims 1,2 page 3, line 5 -page 5, line 7 example 1 figures 1,2	1,4,5, 7-9
A	DE 33 15 105 A (LEUNA WERKE VEB) 17 November 1983 (1983-11-17) the whole document	1,4-9
A	WO 92 05870 A (MONSANTO CO) 16 April 1992 (1992-04-16) claims 1,3,4,10-12 page 16, line 34 -page 17, line 38 page 31; table 1 figures 5A,5B,11A,11B	1-3,7-9
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☒ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

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C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	EP 0 020 963 A (BASF AG) 7 January 1981 (1981-01-07) page 3, line 30 -page 5, line 29 claims 1,2 example ---	1-3,6-9
A	EP 0 004 079 A (HOECHST AG) 19 September 1979 (1979-09-19) claims 1,2 figures 1-6 page 3, line 32 -page 5, line 23 example ---	1-3,5, 7-9
A	EP 0 342 759 A (SHELL INT RESEARCH) 23 November 1989 (1989-11-23) -----	

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/NL 99/00676

Patent document cited in search report		Publication date	Patent family member(s)	Publication date
EP 0008424	A	05-03-1980	DE 2837018 A	06-03-1980
DE 3315105	A	17-11-1983	DD 218736 A	13-02-1985
			BG 47826 A	15-10-1990
			SU 1321460 A	07-07-1987
WO 9205870	A	16-04-1992	AT 125467 T	15-08-1995
			AU 649748 B	02-06-1994
			AU 8859591 A	28-04-1992
			CA 2091767 A	05-04-1992
			CN 1061352 A,B	27-05-1992
			CZ 9300551 A	19-01-1994
			DE 69111612 D	31-08-1995
			DE 69111612 T	25-01-1996
			EP 0552287 A	28-07-1993
			HR 920509 A	31-08-1994
			JP 6501645 T	24-02-1994
			MX 9101444 A	05-06-1992
			NZ 240082 A	26-05-1993
			US 5168090 A	01-12-1992
EP 0020963	A	07-01-1981	DE 2922116 A	11-12-1980
			DK 233280 A	01-12-1980
EP 0004079	A	19-09-1979	DE 2811115 A	27-09-1979
			CA 1122959 A	04-05-1982
			JP 54128490 A	05-10-1979
			US 4370261 A	25-01-1983
			US 4370492 A	25-01-1983
EP 0342759	A	23-11-1989	AU 614136 B	22-08-1991
			AU 3488089 A	23-11-1989
			CA 1335809 A	06-06-1995
			CN 1038033 A	20-12-1989
			DE 68909106 D	21-10-1993
			DE 68909106 T	13-01-1994
			ES 2044055 T	01-01-1994
			FI 892365 A	20-11-1989
			IN 173763 A	09-07-1994
			JP 2026642 A	29-01-1990
			NZ 229161 A	26-04-1991